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Amendments to the Claims

This listing of claims will replace all prior versions and listings of claims in the application:

- 1. (Currently Amended) A method for inhibiting the immunological rejection of a <u>an allogenic</u> transplant in a subject which comprises administering to the subject, at a suitable time <u>after transplant</u>, a <u>prophylactically effective an</u> amount of streptavidin <u>effective to inhibit such immunological rejection</u>.
- 2. (Original) The method of claim 1, wherein the subject is a human.
- 3. (Original) The method of claim 1, wherein the transplant is an organ transplant.
- 4. (Original) The method of claim 1, wherein the transplant is a tissue transplant.
- 5. (Canceled)
- 6. (Canceled)
- 7. (Original) The method of claim 1, wherein the streptavidin is administered intraperitoneally.

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- 8. (Original) The method of claim 7, wherein the streptavidin is administered in a dose of between about 2 mg/kg to about 200 mg/kg of subject body weight per day.
- 9. (Original) The method of claim 8, wherein the streptavidin is administered in a dose of between about 10 mg/kg to about 40 mg/kg of subject body weight per day.
- 10. (Original) The method of claim 9, wherein the streptavidin is administered in a dose of about 20 mg/kg of subject body weight per day.
- 11. (Original) The method of claim 1, wherein the streptavidin is administered intravenously.
- 12. (Original) The method of claim 11, wherein the streptavidin is administered in a dose of between about 2 mg/kg to about 200 mg/kg of subject body weight per day.
- 13. (Original) The method of claim 12, wherein the streptavidin is administered in a dose of between about 10 mg/kg to about 40 mg/kg of subject body weight per day.
- 14. (Original) The method of claim 13, wherein the streptavidin is administered in a dose of about 20 mg/kg of subject body weight per day.
- 15. (Original) The method of claim 1, wherein the streptavidin is administered subcutaneously.

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- 16. (Original) The method of claim 15, wherein the streptavidin is administered in a dose of between about 2 mg/kg to about 200 mg/kg of subject body weight per day.
- 17. (Original) The method of claim 16, wherein the streptavidin is administered in a dose of between about 10 mg/kg to about 40 mg/kg of subject body weight per day.
- 18. (Original) The method of claim 17, wherein the streptavidin is administered in a dose of about 20 mg/kg of subject body weight per day.
- 19. (Original) The method of claim 1, further comprising the step of administering an anti-lymphocyte antibody to the subject at a suitable time.
- 20. (Original) The method of claim 19, wherein the anti-lymphocyte antibody is administered to the subject concurrently with streptavidin.
- 21. (Original) The method of claim 19, wherein the anti-lymphocyte antibody is administered to the subject at a time different from that when streptavidin is administered.

22-26. (Canceled)